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Prolonged Duration of Early Antibiotic Therapy in Extremely Premature Infants

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Abstract

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Dr. Cotten assisted with acquisition of the data, interpreted the data analyses, reviewed and revised the manuscript for important intellectual content, and obtained funding to support the study.

Dr. Stoll assisted with acquisition of data and critical revision of the manuscript for important intellectual content.

Dr. Higgins conceptualized and designed the study, assisted with data analysis and interpretation, and provided critical revision of the manuscript for intellectual content.

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All authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.

Access to data statement: Ms. Chowdhury, Ms. Hansen, and Dr. Das had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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Background: Prolonged early antibiotics in extremely premature infants may have negative effects. We aimed to assess prevalence and outcomes of provision of prolonged early antibiotics to extremely premature infants in the absence of culture-confirmed infection or NEC.

Methods: Cohort study of infants from 13 centers born without a major birth defect from 2008-2014 who were 401-1000 grams birth weight, 22-28 weeks gestation, and survived ≥ 5 days without culture-confirmed infection, NEC, or spontaneous intestinal perforation. We determined the proportion of infants who received prolonged early antibiotics, defined as ≥ 5 days of antibiotic therapy started at ≥ 72 hours of age, by center and over time. Associations between prolonged early antibiotics and adverse outcomes were assessed using multivariable logistic regression.

Results: A total of 5730 infants were included. The proportion of infants receiving prolonged early antibiotics varied from 30-69% among centers and declined from 49% in 2008 to 35% in 2014. Prolonged early antibiotics was not significantly associated with death (adjusted odds ratio 1.17 [95% CI, 0.99-1.40], $p=0.07$) and was not associated with NEC.

Conclusions: The proportion of extremely premature infants receiving prolonged early antibiotics decreased, but significant center variation persists. Prolonged early antibiotics was not significantly associated with increased odds of death or NEC.

INTRODUCTION

Antibiotics are the most common medications prescribed for infants in the neonatal intensive care unit (NICU) (1). Although the incidence of culture-confirmed early-onset sepsis (EOS) in term infants is relatively low (~ 0.5 - 0.8 cases/1000 live births), both the incidence of disease (15-19 cases/1000) and EOS-attributable mortality (3.5% vs 35%) are substantially higher among very low birth weight, preterm infants (2). Given the high mortality associated with EOS, presence of maternal and neonatal risk factors, and time required for pathogens to be isolated in culture, empirical antibiotics targeting EOS are administered to large proportions of infants in the NICU (3,4).

In 2009, the *Eunice Kennedy Shriver* National Institute of Child Health and Human Development Neonatal Research Network (NRN) reported data from 1998-2001 suggesting that prolonged duration of antibiotics administered soon after birth to extremely low birth weight (ELBW, birth weight <1000 g) infants was associated with increased risk of death, necrotizing enterocolitis (NEC), and the composite measure of NEC or death (5). In that study, 53% of 4039 infants surviving >5 days were administered antibiotics beginning at birth and extending ≥ 5 days in the absence of culture-confirmed infection. These findings were consistent with other smaller studies (6,7). It is unclear to what degree neonatal clinicians have changed antibiotic practice over the ensuing 20 years and the impact on infant outcomes. A recent study including 40,534 very low birth weight (VLBW, <1500 g) infants born at 297 primarily nonteaching centers found a small but significant decrease in the rate of prolonged early antibiotic administration among VLBW infants, but no change among the subgroup of 14,923 ELBW infants (8). The purpose of this study was to 1) describe changes over time in the provision of prolonged early antibiotic therapy without culture-confirmed infection or intraabdominal inflammatory process, and 2) describe associations of such prolonged early antibiotic therapy with adverse outcomes among a more

contemporary cohort of extremely premature infants cared for at academic centers of the NRN.

MATERIALS AND METHODS

Subjects

This was a retrospective analysis of data entered into a registry of high-risk preterm infants maintained by the NRN. Study infants were born between January 1, 2008, and December 31, 2014. The registry contains maternal and neonatal information from the intrapartum period until infant discharge home, transfer from the primary center, death, or 120 days after birth. Limited demographic, clinical, and outcome data, including whether or not infants were started on antibiotics before and continued beyond the first 5 postnatal days, were collected by trained research staff utilizing a standardized manual of operations. Site participation differed over the study period, and we included data from the 13 centers that participated in the NRN for the entire study period (denoted with an asterisk in the Acknowledgments). Infants were included in the study if they had a birth weight of 401-1000 g and gestational age of 22 0/7 – 28 6/7 weeks. Exclusion criteria were: death <5 days after birth; blood or cerebrospinal fluid culture obtained within the first 5 postnatal days growing pathogenic bacterial or fungal species; NEC or spontaneous intestinal perforation (SIP) diagnosed ≤ 5 days after birth; or major birth defect. The institutional review board at each center approved participation in the registry.

Definitions

Prolonged early antibiotic therapy was defined as ≥ 5 days of antibiotic administration initiated at ≥ 72 hours of age. Comparator infants may have had antibiotics initiated within 72 hours of age and discontinued before 5 days, or may not have had antibiotics initiated within 72 hours of age. Small for gestational age (SGA) was defined as birth weight <10th percentile for sex and gestational age using percentiles reported by Alexander et al (9). Late-onset sepsis (LOS) was defined as a blood and/or cerebrospinal fluid culture obtained at >72 hours of age and growing a recognized bacterial or fungal pathogen if the infant was administered antibiotics for ≥ 5 days or until death. NEC was defined as modified Bell's stage IIA (10,11).

Statistical analysis

Prior to excluding infants who died at <5 days or who had a positive culture, NEC, or SIP in the first 5 days, we determined the proportion of infants with EOS by center. After exclusions, we determined the proportion of infants who received prolonged early antibiotics overall and by center over time. We compared infant demographic and maternal characteristics between infants who received prolonged early antibiotics and those who did not using Student's t-test for continuous variables and Wald chi-square test for categorical variables. P-values were 2-tailed with values ≤ 0.05 considered statistically significant. Analyses were conducted using SAS 9.4 (SAS Institute, Cary, NC).

Separate logistic regression models were used to evaluate associations between birth year, center, and receipt of prolonged early antibiotics; and between prolonged early antibiotics

and outcomes, including all-cause mortality after 5 days of life, NEC, LOS, LOS due to organisms other than coagulase-negative staphylococci (non-CoNS LOS), and LOS due to one or more fungal organisms (fungal LOS). We examined death and composite outcomes of death/morbidity for all infants; NEC and LOS were examined in survivors to discharge to reduce potential bias due to different at-risk periods, particularly for infants with short survival times. All models included birth year (categorical), study center, infant characteristics (gestational age, SGA, sex, and 5-minute Apgar score <5), and maternal characteristics (black race, multiple birth, rupture of membranes >24 hours, receipt of antenatal steroids, perinatal antibiotic therapy, hypertension, antepartum hemorrhage, and cesarean section).

Because prolonged early antibiotics may be administered to more severely ill infants at greater risk for adverse outcomes, we also included an indicator in our models aimed at identifying severity of illness. We defined this indicator as initiation of any mechanical ventilation on the day of birth and continued use for all of the first 7 days after birth, or for all days until death if death occurred before 7 days (continuous mechanical ventilation [cMV]). We examined whether associations between prolonged early antibiotics and adverse outcomes were different among infants with and without this indicator by including the interaction between prolonged early antibiotics and cMV in our models. Results are shown separately for the cMV and non-cMV groups if this interaction was significant at $p \leq 0.10$. Otherwise, results shown are based on models that did not include the cMV interaction.

RESULTS

The NRN registry enrolled 14,212 infants from 2008-2014. Of these, 5730 met all study inclusion and exclusion criteria (Supplemental Digital Content S1). Prolonged early antibiotics were administered to 2526/5730 (44%) infants. Multiple maternal and neonatal characteristics differed significantly between infants who did or did not receive prolonged early antibiotics (Table 1). Overall, infants who received prolonged early antibiotics were younger and smaller, had lower 5-minute Apgar scores, were less likely to have been born via cesarean section, were more likely to have been born to a mother with prolonged rupture of membranes or who was treated with antenatal antibiotics, and were more likely treated with cMV (Table 1).

Factors associated with receipt of prolonged early antibiotics

The overall percentage of infants treated with prolonged early antibiotics decreased from 49% in 2008 to 35% in 2014 (adjusted odds ratio [aOR] 0.44 [95% CI, 0.35-0.56], $p < 0.001$) (Supplemental Digital Content S2, Table 2). The percentage of infants treated varied by center, ranging from 30-69% (Figure 1). The center-specific incidence of EOS ranged from 0.3-3.6%. The center-specific rate of prolonged early antibiotic administration decreased over the study period at 8 of the 13 centers. Compared to the center with the lowest rate, infants cared for at 11 of the 12 remaining centers were significantly more likely to receive prolonged early antibiotics (Table 2).

Relationship of prolonged early antibiotics with outcomes

Infants who received prolonged early antibiotics had marginally but not significantly higher odds of death compared to those who did not receive prolonged early antibiotics (19% vs 13%, aOR 1.17 [95% CI, 0.99-1.40], $p=0.07$). Prolonged early antibiotics were not significantly associated with the composite outcomes of death or specific morbidities (Table 3), nor with specific morbidities among survivors to discharge. These findings were not different among infants who did or did not receive cMV, with one exception: the relationship of prolonged early antibiotics with LOS caused by fungal organisms varied by cMV group (interaction $p<0.001$) (Table 3). Among infants not treated with cMV, prolonged early antibiotic administration was associated with increased odds of fungal LOS (2% vs 0.4%, aOR 4.95 [95% CI, 1.93-12.69], $p<0.001$) (Table 3).

DISCUSSION

We observed a significant decline in the overall use of prolonged early antibiotics among ELBW infants cared for in 13 NRN centers over a 7-year period. The overall use of prolonged early antibiotic therapy during this study was lower (44% vs 53%) than that reported in our prior study of early antibiotic use in infants born 1998-2001. However, we observed persistent significant center-specific antibiotic practice variation. The proportion of infants treated with prolonged early antibiotics differed between centers, and reduction in prolonged therapy was only observed at 8 of the 13 centers. The rates of prolonged antibiotic administration remained 10-to-100-fold higher than the rates of culture-confirmed EOS, even with the observed decrease in prolonged therapy. In contrast to our prior NRN study, we found no statistically significant association between prolonged early antibiotics and risk of death. Our study raises several issues to be considered as neonatal caregivers work to identify the best approaches to use of antibiotics for medically fragile, extremely premature infants in the first postnatal days.

Our findings reflect slowly evolving views of the role of antibiotics in neonatal care. Multiple national organizations recommend comprehensive approaches to antimicrobial stewardship (12), and neonatal providers are actively pursuing approaches to optimize antibiotic use among NICU patients (13). In a study using the Pediatric Health Information System database, antibiotic utilization decreased ~5% per year from 2004 to 2012 among hospitals without mandatory public reporting requirements for central line-associated bloodstream infection (14); similar secular trends in NICU antibiotic use were observed among hospitals subject to such reporting requirements. Neonatal antimicrobial stewardship programs at individual centers and collaboratives also report success in safely decreasing overall NICU antibiotic utilization (15-17). However, NICUs care for infants across a wide spectrum of gestational ages and morbidities, and studies focused specifically on VLBW and ELBW infants report mixed outcomes. The Premier Perspective Database study measured quarterly rates of antibiotic initiation and duration among 40,364 VLBW infants cared for in 297 centers across the United States from 2009-2015 (8). No significant trends were observed for antibiotic initiation, but declines were noted for prolonged early antibiotics among VLBW (but not ELBW) infants. The Canadian Neonatal Network (CNN) reported decreased antibiotic use in VLBW infants between 2010 and 2014, a finding that coincided

with decreased rates of LOS (18). Despite this decline, the study still found an overall 20% antibiotic utilization rate among VLBW infants in the absence of culture-confirmed infection or NEC. Single-site studies further illustrate the challenges of antibiotic optimization among premature infants. Guidelines for use of antibiotics targeting highly antibiotic-resistant pathogens (e.g., meropenem, vancomycin) resulted in more appropriate antibiotic use in one Canadian NICU, but improvements were not seen among VLBW infants (19). A comprehensive antibiotic stewardship program in a large American NICU did not significantly decrease overall antibiotic utilization, although some changes were observed in the use of specific antibiotics such as ampicillin and vancomycin (20). Viewed from the perspective of the many challenges in optimizing neonatal antibiotic administration, the decline in prolonged early antibiotic administration we observed in the current study (14% absolute change, 28% relative change) among ELBW infants without a corresponding statistically significant rise in subsequent mortality, LOS, or NEC is clinically significant.

One important issue raised by our study is whether there is a critical level, duration, or timing of antibiotic exposure at which antibiotics administered in the absence of proven infection begin to cause harm. In contrast to prior findings from the NRN and other centers (6,21), we found no significant association between prolonged early antibiotics and NEC. Multiple practice differences over the past 20 years may have contributed to this finding, including changes in the use of mothers' own milk, human donor milk, and probiotic supplements, and we did not collect detailed information on those factors. Our findings do suggest, however, that the association of empiric antibiotic use and risk of NEC has declined as overall antibiotic use has declined. The multicenter CNN study also suggests that the magnitude of antibiotic exposure may be associated with the magnitude of antibiotic-related harm (18). Increased exposure to antibiotics over an infant's entire hospital stay was associated with increased adjusted odds of a composite outcome of mortality and major morbidities, with the greatest risk associated with the highest quartile of antibiotic exposure. The prior NRN study noted increased odds of death per day of antibiotic therapy (OR=1.16 [95% CI, 1.08-1.24]), and a substantial proportion of infants received therapy for 10 days (5). In the current study, neither duration of antibiotics nor daily antibiotic exposure data were recorded, and it is possible that our findings reflect an even larger magnitude of decreased antibiotic use than we have measured.

We continue to find substantial center variation in the use of prolonged early antibiotics (range of 30-69% from 2008-2014 compared to 27-85% in 1998-2001). This variation appeared to be unrelated to the incidence of EOS at each site. It is unlikely to be due to a corresponding variation in severity of illness, as each NRN site is an academic, tertiary care center caring for relatively large numbers of extremely premature infants. Schulman and colleagues also found wide variation in California NICU antibiotic use unexplained by variation in proven infection, NEC, surgical volume, or mortality (22). We speculate that center variation in this early postnatal clinical practice represents true elective practice differences that reflect provider uncertainty regarding the risks and benefits of antibiotics administered in the absence of proven infection. In our cohort, simply being born at one academic center instead of another could increase an infant's odds of receiving prolonged early antibiotics sevenfold, even after adjustment for demographic and clinical factors. Our

findings underscore an imperative for individual centers to track antibiotic utilization and to compare such data with other institutions. Robust mechanisms for sharing and receiving such data, including membership in voluntary neonatal networks, will facilitate efforts to improve local practices.

Potentially unjustified variation may matter. While not statistically significant, receipt of prolonged early antibiotics was associated with marginally increased odds of death in our cohort of infants (aOR 1.17 [95% CI, 0.99-1.40]). An OR >1.0 is consistent with the previous NRN report, which showed an aOR of 1.46 (95% CI, 1.19-1.78) for prolonged therapy (5). Although our analysis cannot fully account for differences in severity of illness among infants who do and do not receive prolonged early antibiotics, the potential for mediating harm remains of concern. The association between prolonged early antibiotics and fungal LOS in survivors to discharge may be an example of harm rather than confounding by indication. In this instance, relative harm was associated with prolonged early antibiotic administration to infants who were clearly *less* severely ill (those whose caregivers did not decide to use cMV) compared to those who were more severely ill (those who were treated with cMV). Broad-spectrum antibiotic use has been previously associated with invasive candidiasis in premature infants (23,24), although why such use might differentially affect those of less illness severity, defined by available data describing degree of respiratory support, remains unclear. We speculate that less severely ill infants had fewer risk factors for fungal LOS and thus risk was increased significantly due to the use of prolonged antibiotics; more severely ill infants may have already had maximized risk of fungal LOS due to other factors. Increased perinatal and postnatal exposure to antibiotics in premature infants has been associated with alterations in the gut microbiome, and may provide some insight into the biologic mechanisms by which prolonged antibiotic administration may be harmful in the absence of confirmed infection (25,26). It is possible that dysbiosis leads to proliferation of potential pathogens and predisposition to later infection, with resulting increases in morbidity and mortality (27). In addition, antibiotic exposure has been associated with a reduction in gastric bacterial colonization, which could increase risk for infection and increase feeding intolerance (28). Regardless of the potential mechanisms of harm or magnitude of harm, our findings support ongoing efforts to limit empiric antibiotic administration to ELBW infants at highest risk of infection and to spare those who may be at lowest risk of EOS, such as those born by cesarean section, without labor or maternal chorioamnionitis (29).

The strengths of our study include the large sample size of extremely preterm infants, uniform definitions and data collection over time, and observation of trends over time and across multiple regions in the United States. The retrospective analysis limits our ability to measure variables other than those routinely collected in the NRN database. We attempted to account for clinical severity by adjusting for mechanical ventilation, but we were unable to calculate validated severity scores such as the CRIB II or SNAPPE-II (30,31). While we excluded infants with culture-confirmed sepsis in the first 5 postnatal days, we were unable to identify infants who may have had other true infections, such as neonatal pneumonia. We did not have data regarding the type or timing of quality improvement interventions or antibiotic stewardship programs at participating centers. Finally, while our study was similar to the previous NRN study, several differences limit comparisons between the two reports.

The previous study included infants diagnosed with NEC or SIP in the first 5 days after birth, while we excluded these infants (Supplemental Digital Content S1). Our rationale was that antibiotic administration to those infants is not discretionary, and infants with NEC and SIP are likely to justifiably receive prolonged early antibiotics and have increased likelihood of mortality and morbidity. The previous NRN study included expanded data collection that detailed the exact days of exposure and the type of antibiotics prescribed on each day. This level of detail was not available to the current study. In the earlier NRN study, even 4 days of early antibiotics was associated with NEC, death, as well as the composite outcome; but in the current study, we were unable to examine the effect of antibiotics administered for >48 hours but <5 days, and the risk of this practice remains unclear. We also do not know what proportion of infants had no antibiotics given in the first 72 hours after birth (~2-3% had none given in the prior study) (5), and we do not know what proportion of infants were administered cephalosporins, antibiotics that are associated with higher mortality and *Candida* infections (24,32).

CONCLUSION

In this study of 5730 infants across 13 NRN centers, the proportion of infants who received prolonged early antibiotics in the absence of EOS, NEC, or SIP decreased significantly between 2008 and 2014. Overall prolonged antibiotic use also declined in comparison to a prior NRN study that included infants born between 1998 and 2001. Prolonged early antibiotic exposure continues to vary widely across NRN centers. Our findings support ongoing efforts to promote neonatal antibiotic stewardship, as supported by the Centers for Disease Control and Prevention (33). Such efforts must include robust means of identifying those infants who will derive maximum benefit and minimum harm from antibiotic treatments that will be acceptable across different sites of neonatal care.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Data collected at participating sites of the NICHD Neonatal Research Network (NRN) were transmitted to RTI International, the data coordinating center (DCC) for the network, which stored, managed, and analyzed the data for this study. On behalf of the NRN, Dr. Abhik Das (DCC Principal Investigator) and Ms. Dhuly Chowdhury and Ms. Nellie Hansen (DCC Statisticians) had full access to all the data in the study and take responsibility for the integrity of the data and accuracy of the data analysis. The sites with data included in this study are denoted with an asterisk.

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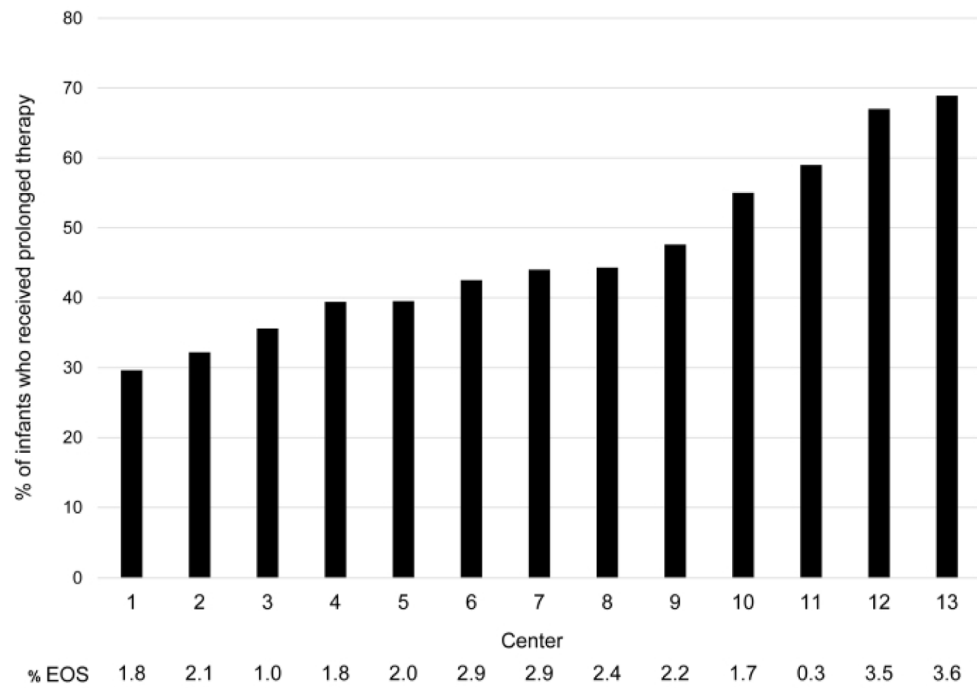


Figure 1. Percentage of study infants receiving prolonged early antibiotic therapy and percentage of infants with early-onset sepsis (EOS) by center.

Table 1.Demographics and clinical characteristics^a

	Prolonged early antibiotics (N=2526)	No prolonged early antibiotics (N=3204)	P-value
Infant characteristics			
Birth weight, g (median, IQR)	730 (615, 850)	790 (670, 890)	<0.001
Gestational age, weeks (median, IQR)	25 (24, 26)	26 (25, 27)	<0.001
By gestational age week			<0.001
22	1%	1%	
23	11%	6%	
24	23%	15%	
25	24%	21%	
26	20%	23%	
27	13%	21%	
28	7%	14%	
Small for gestational age	11%	11%	0.48
Male	51%	46%	<0.001
5-minute Apgar <5	24%	17%	<0.001
Mechanically ventilated for first 7 days of life ^b	55%	33%	<0.001
Maternal characteristics			
Race/ethnicity ^c			0.19
Black	45%	43%	
White	36%	38%	
Hispanic	13%	14%	
Other	6%	5%	
Multiple birth	23%	26%	0.01
Hypertension	26%	37%	<0.001
Antenatal steroids	89%	91%	0.05
Antenatal antibiotics	76%	71%	<0.001
Rupture of membranes >24 hours	28%	18%	<0.001
Antepartum hemorrhage	19%	18%	0.84
Cesarean section	64%	73%	<0.001

IQR=interquartile range

^aInformation was missing as follows: maternal race/ethnicity, 16 infants; maternal hypertension, 8 infants; antepartum hemorrhage, 5 infants; antenatal steroids, 12 infants; antenatal antibiotics, 34 infants; timing of rupture of membranes, 159 infants; cesarean delivery, 3 infants; 5-minute Apgar score, 14 infants; mechanical ventilation, 1 infant.

^bThis group includes 40 infants who were intubated continuously until death before day 7.

^cMaternal race/ethnicity was defined as non-Hispanic white, non-Hispanic black, Hispanic (both white and black), and other race (Asian/Pacific Islander, American Indian / Alaska native, more than one race, other not specified). Those of white or black race with missing ethnicity information (5.3% of blacks, 0.9% of whites) were classified as non-Hispanic.

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Table 2.

Differences in receipt of prolonged early antibiotic therapy by birth year and center adjusting for neonatal and maternal characteristics

	Adjusted odds ratio (95% confidence interval)	P-value
Birth year (reference: 2008)		
2009	0.89 (0.72-1.10)	0.29
2010	1.02 (0.82-1.27)	0.83
2011	0.79 (0.63-0.98)	0.03
2012	0.58 (0.47-0.72)	<0.001
2013	0.58 (0.47-0.73)	<0.001
2014	0.44 (0.35-0.56)	<0.001
Center (reference: Center 1)		
2	1.16 (0.90-1.50)	0.26
3	1.42 (1.10-1.83)	0.008
4	1.78 (1.33-2.39)	<0.001
5	1.61 (1.14-2.27)	0.007
6	2.24 (1.53-3.28)	<0.001
7	1.61 (1.20-2.16)	0.002
8	1.81 (1.34-2.43)	<0.001
9	2.17 (1.67-2.82)	<0.001
10	2.97 (2.20-4.01)	<0.001
11	3.07 (2.26-4.17)	<0.001
12	5.46 (4.03-7.39)	<0.001
13	7.36 (5.18-10.46)	<0.001
Infant characteristics		
Gestational age, weeks (reference: 28)		
22	1.81 (0.96-3.41)	0.07
23	2.54 (1.86-3.47)	<0.001
24	1.98 (1.54-2.55)	<0.001
25	1.74 (1.38-2.20)	<0.001
26	1.40 (1.11-1.76)	0.004
27	1.06 (0.83-1.34)	0.66
Small for gestational age	1.65 (1.35-2.01)	<0.001
Male	1.15 (1.03-1.29)	0.02
5-minute Apgar <5	1.38 (1.19-1.61)	<0.001
Mechanically ventilated for entire first 7 days of life	1.79 (1.57-2.05)	<0.001
Maternal characteristics		
Black race	1.24 (1.08-1.41)	0.002
Multiple birth	0.90 (0.78-1.04)	0.16

	Adjusted odds ratio (95% confidence interval)	P-value
Hypertension	0.78 (0.67-0.91)	0.001
Antenatal steroids	0.82 (0.66-1.02)	0.07
Antenatal antibiotics	1.18 (1.01-1.37)	0.03
Rupture of membranes >24 hours	1.84 (1.59-2.14)	<0.001
Antepartum hemorrhage	0.83 (0.71-0.97)	0.02
Cesarean section	0.84 (0.73-0.97)	0.02

Table 3.

Outcomes for infants who received prolonged early antibiotic therapy versus those who did not

Outcome, n (column %)	Prolonged early antibiotics	No prolonged early antibiotics	Adjusted OR for outcome (95% CI) prolonged early antibiotics vs no prolonged early antibiotics	P-value
All infants who survived 5 days	2526	3204		
Death before discharge	466 (19)	409 (13)	1.17 (0.99-1.40)	0.07
Death or NEC	623 (25)	635 (20)	1.08 (0.93-1.25)	0.34
Death or LOS	1008 (40)	1082 (34)	0.98 (0.86-1.12)	0.76
Death or non-CoNS LOS	662 (26)	677 (21)	1.01 (0.87-1.17)	0.89
Death, NEC, or LOS	1086 (43)	1204 (38)	0.97 (0.86-1.11)	0.66
Infants who survived to discharge	2055	2787		
NEC	157 (8)	225 (8)	0.91 (0.72-1.15)	0.43
LOS	540 (26)	670 (24)	0.90 (0.77-1.05)	0.18
Non-CoNS LOS	195 (10)	268 (10)	0.84 (0.67-1.04)	0.11
Fungal LOS ^a				
cMV group	23 (2)	22 (3)	0.64 (0.33-1.23)	0.18
Non-cMV group	21 (2)	8 (0.4)	4.95 (1.93-12.69)	<0.001

OR=odds ratio; CI=confidence interval; NEC=necrotizing enterocolitis; LOS=late-onset sepsis; CoNS=Coagulase-negative *Staphylococcus*; cMV=continuous mechanical ventilation for first 7 days.

^aThe interaction between prolonged early antibiotics and the continuous mechanical ventilation group was significant at $p<0.001$, and therefore the adjusted ORs are reported separately.